

PATENT

Docket No.: 201040/1020

Examiner: H. Schnizer

Art Unit:

1653

TROHOENTER TO TOO STORE TO

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant(s) :

Alam et al.

Serial No.

09/455,978

Cnfrm. No.

5811

Filed

December 6, 1999

For

HEME PROTEINS HEMAT-HS AND

HEMAT-BS AND THEIR USE IN

MEDICINE AND MICROSENSORS

U.S. Patent and Trademark Office

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Dear Sir:

Transmitted herewith in the above-identified application are:

- Statement in Accordance with 37 C.F.R. § 1.821(g) and a substitute computer [X]readable 3.5" Diskette as required by 37 C.F.R. § 1.825(d).
- XCopy of Notice to Comply with Requirements for Patent Applications Containing Nucleotide Sequence and/or Amino Acid Sequence Disclosures; and
- [X]A self-addressed postcard for acknowledging receipt.

[X]The Commissioner is hereby authorized to charge any additional fees or credit any overpayment to Deposit Account No. 14-1138 .

A duplicate copy of this sheet is enclosed

Date: March 11, 2002

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Application No.: <u>09/455,978</u> NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Applicant must file the items indicated below within the time period set the Office action to which the Notice is attached to avoid abandonment under 35 U.S.C. § 133 (extensions of time may be obtained under the provisions of 37 CFR 1.136(a)).

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

	1.	This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to the final rulemaking notice published at 55 FR 18230 (May 1, 1990), and 1114 OG 29 (May 15, 1990). If the effective filing date is on or after July 1, 1998, see the final rulemaking notice published at 63 FR 29620 (June 1, 1998) and 1211 OG 82 (June 23, 1998).
	2.	This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c).
	3.	A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).
	4.	A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing."
X	5.	The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).
	6.	The paper copy of the "Sequence Listing" is not the same as the computer readable from of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).
	7.	Other: _
	Applicant Must Provide:	
X	An	initial or substitute computer readable form (CRF) copy of the "Sequence Listing".
	An initial or substitute paper copy of the "Sequence Listing", as well as an amendment directing its entry into the specification.	
X	A s	statement that the content of the paper and computer readable copies are the same and, where applicable, lude no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g).

For questions regarding compliance to these requirements, please contact:

- For Rules Interpretation, call (703) 308-4216 or (703) 308-2923
- For CRF Submission Help, call (703) 308-4212
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